



A new golden frog species of the genus Diasporus (Amphibia, Eleutherodactylidae) from the Cordillera Central, western Panama

Andreas Hertz^{1,2,†}, Frank Hauenschild^{1,2,‡}, Sebastian Lotzkat^{1,2,§}, Gunther Köhler^{1,1}

I Senckenberg Forschungsinstitut und Naturmuseum, Senckenberganlage 25, 60325 Frankfurt am Main, Germany **2** Johann Wolfgang Goethe-University, Institute for Ecology, Evolution and Diversity, Biologicum, Building C, Max-von-Laue-Straße 13, 60438 Frankfurt am Main, Germany

- † urn:lsid:zoobank.org:author:A22AE5AB-B67A-4140-9CF5-F7DD1FAA7DE5
- ‡ urn:lsid:zoobank.org:author:4B072D05-5B17-44C1-AE6D-137B5BBB5856
- § urn:lsid:zoobank.org:author:886EF473-1B7B-4EA7-8142-FCD42CEEF903
- urn:lsid:zoobank.org:author:71305C99-4BD6-4305-A884-9F8221EBA11B

Corresponding author: Andreas Hertz (ahertz@senckenberg.de)

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Abstract

We describe the frog species *Diasporus citrinobapheus* **sp. n.** from the Cordillera Central of western Panama. The new species differs from all other species in its genus in coloration, disk cover and disk pad shape, skin texture, advertisement call, and size. It is most similar to *D. tigrillo*, from which it differs in dorsal skin texture, relative tibia length, number of vomerine teeth, ventral coloration, dorsal markings, and relative tympanum size, and to *D. gularis*, from which it can be distinguished by the lack of membranes between the toes, adult size, posterior thigh coloration, and position of the choanae. We provide data on morphology, vocalization, and distribution of the new species, as well as brief information on its natural history.

Resumen

Describimos la especie de rana *Diasporus citrinobapheus* **sp. n.** de la Cordillera Central, occidente de Panamá. La nueva especie se distingue de otras especies del género por su coloración, su forma de la cubierta y la almohadilla de los discos, textura de la piel, canto de anúncio, y tamaño corporal. Se asemeja mas a *D. tigrillo*, del cual se distingue por la textura de la piel dorsal, longitud relativa de la tibia, número de dientes vomerianos, coloración ventral, patrón dorsal, y tamaño relativo del tímpano, y a *D. gularis*, del

cual se diferencia por la ausencia de membranas entre los dedos de pie, tamaño corporal, coloración de la parte trasera del muslo, y posición de las coanas. Presentamos datos de la morfología, vocalización, y distribución de la nueva especie, así como notas concisas de su historia natural.

Keywords

Central America, Anura, diversity of species, taxonomy, vocalization

Palabras clave

América Central, Anura, diversidad de especies, taxonomía, vocalización

Introduction

Panama's herpetofauna is known to be the most diverse in consideration of its size in Central America, with only Mexico being more diverse in absolute species count (Myers and Duellman 1982; Jaramillo et al. 2010). Although herpetological research has been conducted in Panama for more than a hundred years (Ibáñez et al. 2001), the knowledge of amphibian species diversity is still far from being completed. This is demonstrated impressively by the multitude of amphibian species described from this country within the last years (e.g. Wake et al. 2005, 2007; Köhler et al. 2007; Mendelson et al. 2008; Bolaños and Wake 2009; Crawford et al. 2010b; Mendelson and Mulcahy 2010; Ryan et al. 2010a, 2010b).

The genus *Diasporus* (Hedges et al. 2008) is the closest relative of the predominantly Caribbean genus *Eleutherodactylus*. The species of *Diasporus* are distributed from eastern Honduras to western Ecuador (Frost 2011; Köhler 2011). The genus contains nine described species, five of which (*Diasporus diastema* Cope, *D. hylaeformis* Cope, *D. tigrillo* Savage, *D. ventrimaculatus* Chaves, García-Rodríguez, Mora and Leal, and *D. vocator* Taylor) are currently known to occur in western Panama and/or southern Costa Rica. The remaining four species (*D. anthrax* Lynch, *D. gularis* Boulenger, *D. quidditus* Lynch, and *D. tinker* Lynch) are distributed in Panama east of the Canal, and further along the Pacific coast of northern South America southwards to northwestern Ecuador (Frost 2011; IUCN 2011). However, differences in body size, male advertisement call, and coloration in the genus *Diasporus* suggest that there are several undescribed species (Lynch and Duellman 1997; Ibáñez et al. 1999; Savage 2002). Recent fieldwork in the Serranía de Tabasará of western Panama resulted in the discovery of a remarkable new species of *Diasporus*. The purpose of this paper is to describe this new species.

Materials and methods

Field work was carried out between May and August 2010 at several sites along both slopes of the Serranía de Tabasará between the Fortuna depression and Santa Fé, Veraguas, Panama. All specimens were encountered during opportunistic searches at night.

Preparation and preservation of voucher specimens follows Köhler (2001). Tissue samples, usually the left forearm, were stored in 98% undenatured ethanol and deposited in the tissue collection of the Senckenberg Forschungsinstitut und Naturmuseum Frankfurt, Germany (SMF). Geographic coordinates and altitude above sea level were recorded with a handheld Garmin GPS MAP 60 CSx GPS receiver. All georeferences are in the geographical coordinate system and WGS 1984 datum, given in decimal degrees rounded to the fourth decimal place. Elevations are rounded to the next tenth. The map was created using ArcGIS 10 (ESRI).

We took additional morphological data from all Central American species currently assigned to the genus *Diasporus* in the SMF collection. We list all specimens examined for comparison in the Appendix I. Abbreviations for museum collections follow those of Sabaj Pérez (2010) except MHCH (Museo Herpetológico de Chiriquí, the herpetological collection of the Universidad Autónoma de Chiriquí, David, Panama). Specimens in the Appendix labeled with AH field numbers will be deposited at MHCH.

The sex of the male holotype and the paratypes was determined by the presence of vocal slits and vocal sac. Measurements were made with a dial caliper with the aid of a dissecting microscope and rounded to the nearest 0.1 mm. Measurements of the holotype (LACM 146212) and paratype (LACM 146241) of Diasporus tigrillo were taken from Savage (1997), those of D. ventrimaculatus from Chaves et al. (2009). Additionally, we examined photographs of the type specimens of *D. tigrillo*. If possible, missing measurements were calculated on the basis of data presented in the respective descriptions. In the case of *D. ventrimaculatus*, no individual measurements are available, for which reason calculations were made using the average values given for the paratypes by Chaves et al. (2009). We follow Savage (1997, 2002) in the terminology of disk cover and disk pad shape, dorsal outline of head, and snout profile shape. Abbreviations used for measurements are: snout-vent length (direct line distance from the tip of the snout to the posterior margin of the vent): SVL; length of Finger III (from distal end of the Finger III including disk to the base of the second subarticular tubercle): LF III; length of Toe IV (from distal end of the toe IV including disk to the base of third subarticular tubercle): LT IV; disk width at Finger III (at greatest width): DWF III; disk width at Toe IV (at greatest width): DWT IV; head length (from quadratojugal region to tip of the snout): HL; head width (between angles of jaw): HW; tibia length (straight length of the tibia): TL; horizontal eyelid diameter (greatest length of the upper eyelid): EL; interorbital distance (the width of frontoparietal bone between the orbits): IOD; horizontal tympanum diameter (at greatest length): TD; and horizontal eye diameter (at greatest length): ED. The capitalized colors and color codes (the latter in parentheses) in color descriptions in life are those of Smithe (1975–1981). Drawings of head, hands, and feet were made using a camera lucida attachment for a Leica MZ 12 dissecting microscope. Values provided for morphometric and acoustic parameters are minimum, maximum, and mean value ± standard deviation.

Advertisement calls were recorded using a Sennheiser ME 66 shotgun microphone capsule with a Sennheiser K6 powering module in combination with the Marantz PMD 620 solid-state recorder. A minimum distance from microphone to frog of one

meter was kept while recording to prevent disturbance. As needed, the microphone was attached to branches with the aid of a Joby Gorillapod in order to minimize disturbance of the calling frog. Calls were recorded in PCM format at a sampling rate of 48 kHz with 24 bit resolution and stored as way files on a SD Card. Call editing and analysis were performed using SOUND RULER 0.9.6.0 (Gridi-Papp 2003–2007) for frequency analysis and to generate figures of oscillograms and audiospectrograms. We measured temporal parameters by hand using ADOBE AUDITION 3.0, because SOUND RULER has difficulties in accurately and precisely measuring temporal parameters (Bee 2004). Frequency information was obtained through Fast Fourier Transformation (FFT length 512 points, overlap between FFTs 0.8) at Hanning window function. Air temperature and humidity were measured immediately after each sound recording using the digital device Voltcraft HT-200 to the nearest 1°C and 3.5% relative humidity (RH). An alcohol extraction of skin secretions of the new species has been examined for alkaloids using Liquid Chromatography-Time-Of-Flight-Mass Spectrometry (LC-TOF-MS) at the Center of Forensic Medicine of the Goethe-University Frankfurt am Main.

For the complementary molecular analysis, we extracted DNA following the protocol of Ivanova et al. (2006). To eliminate potential PCR-inhibiting contaminants, the tissue samples were incubated for one hour in TE-buffer (10 parts Tris-HCl (pH = 8.0) and one part EDTA) before digestion for at least 10 hours in 50 µl Vertebrate Lysis Buffer and 5.2µl Proteinase K at 56 °C. After extraction, DNA was eluted in 50 µL TE buffer. A fragment of the mitochondrial 16S rRNA gene was amplified in an MJ Research Dyad Deciple™ Peltier Thermal Cycler using the following program: initial denaturation for 180 s at 94 °C; followed by 39 cycles with denaturation for 15 s at 94 °C, hybridization for 60 s at 51 °C, and elongation for 60 s at 72 °C; final elongation for 120 s at 72 °C. Reaction mix for each sample contained 1 μL DNA template, 2.5 μL Reaction Buffer (PeqGold), 2.5 µL 2.5 mM dNTPs, 0.5 µL Taq Polymerase (PeqLab), 16.5 μL H₂O, and 1 μL of each primer (forward: 16SA-L, 5'-CGCCTGTTTAT-CAAAAACAT-3'; reverse:16SB-H, 5'-CCGGTCTGAACTCAGATCACGT-3'). The achieved 16S sequences were deposited in GenBank. We compared 21 sequences of the genus *Diasporus* in our analysis, three of the type series of the new species and one referred specimen, four specimens referred to as D. aff. hylaeformis from Cerro Pando, as well as all 16S sequences of the genus *Diasporus* available on GenBank. We used an additional 16S sequence of *Pristimantis ridens* as an outgroup (see Appendix II for examined specimens and GenBank accession numbers). Sequences were aligned with ClustalW (Larkin et al. 2007) using the default settings in Geneious (Drummond et al. 2010). The manually refined final alignment contained 535 positions. Using MEGA5 (Tamura et al. 2011), we computed uncorrected pairwise genetic distances, determined the Tamura 3-parameter model (Tamura 1992) as the best-fitting substitution model, and conducted Maximum Likelihood as well as Maximum Parsimony analyses (each with 10000 bootstrap replicates). Using TCSv1.21 (Clement et al. 2000), we conducted a statistical parsimony network analysis, with gaps considered as a fifth character state and a connection limit of 95%.

Results

Diasporus citrinobapheus sp. n.

urn:lsid:zoobank.org:act:4A526693-CA45-44FC-9D9D-4F3064A47341 http://species-id.net/wiki/Diasporus_citrinobapheus Figures 1 A, B; 2; 3; 5

Holotype. Adult male SMF 89814: collected on June 26, 2010 at 19:13 by Andreas Hertz and Sebastian Lotzkat at Quebrada Rasca (8.4851°N, 81.1727°W, 790 m elevation), near Paredón, Comarca Ngöbe-Buglé, western Panama, approximately 50 airline km NNW of the city of Santiago and 20 airline km N of Cañazas, Veraguas.

Paratypes. All collected by Andreas Hertz and Sebastian Lotzkat at the type locality on June 26, 2010: MHCH 2370-71; SMF 89816; all adult males.

Referred specimens. Adult males SMF 89817 and MHCH 2372: collected on July 01, 2010 by Andreas Hertz and Sebastian Lotzkat at the private reserve Willie Mazú, Comarca Ngöbe-Buglé (8.7903°N, 82.1989°W, 681 m elevation); female SMF 89820: collected on March 31, 2009 by Andreas Hertz, Sebastian Lotzkat and Arcadio Carrizo at Cerro Negro, Parque Nacional Santa Fé, Veraguas (8.5691°N, 81.0988°W, 730 m elevation).

Diagnosis. A member of the genus *Diasporus* based on the following combination of characters: vocal slits and a single subgular vocal sac present, adult males without nuptial thumb pads; Finger I shorter than Finger II; Toe III much shorter than Toe V; subarticular tubercles on hands and feet flattened; no supernumerary tubercles on hands and feet; no tarsal fold or tubercle. Diasporus citrinobapheus differs from all described members of its genus by the following combination of characters (for accounts, see Table 1): coloration bright yellow to orange in life (Fig. 1 A); head almost as broad as long, but comparatively broad in relation to SVL; skin of dorsum smooth; venter coarsely areolate; tympanum covered by skin but annulus clearly visible; TD about 41% of ED; EL on average narrower than IOD; snout subacuminate in profile and rounded to subovoid in dorsal outline; disks of fingers and toes slightly expanded, disk covers of most fingers and toes spadate, but lacking papillae; disk pads of most fingers and toes triangular; subarticular tubercles of hands and feet rounded, very flat, almost not visible; vomerine odonthophores longish oval and widely separated; vomerine teeth weakly developed; upper eyelid usually smooth, very low pustules in some individuals; heel smooth. Its bright yellow to orange coloration distinguishes D. citrinobapheus from almost all described Central American Diasporus, which, in spite of considerable variation, are all tan to gray or brownish to almost black. In D. hylaeformis and D. ventrimaculatus, the dorsal ground color can be suffused with pink or red. Only D. tigrillo from Costa Rica, a species known only from two specimens, shows a yellowish coloration in life according to the original description (Savage 1997). Diasporus citrinobapheus differs from the two known specimens of D. tigrillo in the following characters (character for *D. tigrillo* in parentheses): SVL in adult males 17.3–19.7 mm (16.0–17.5 mm); dorsal skin absolutely smooth (dorsal skin with scattered low pus-

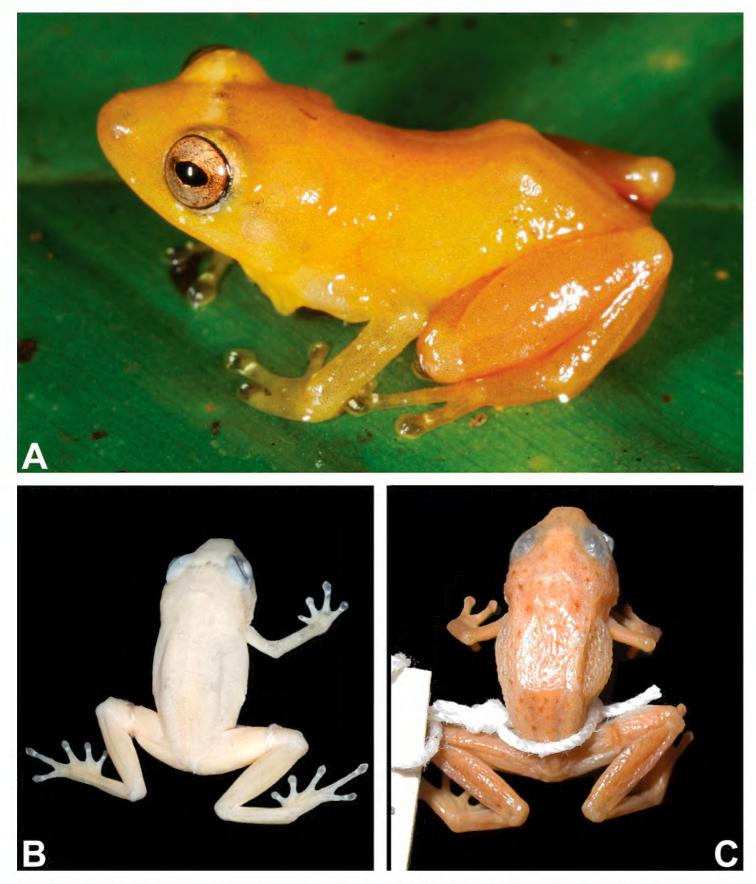


Figure 1. A–B Holotype of *Diasporus citrinobapheus* (SMF 89814, adult male): **A** in life **B** in preservative. **C** *Diasporus tigrillo* in preservative (LACM 146212, holotype, adult male), note dark brown spots. Pictures are not at the same scale.

tules, best developed on dorsum); TD 32–45% of ED (54–57%); TL 40% of SVL (about 48%); distal subarticular tubercle of Finger and Toe I flat and rounded (weakly bifid); many weakly developed vomerine teeth in three to four close rows (a few vomerine teeth in two obliquely aligned and widely separated rows); dorsal surface uniformly bright yellow to orange, sometimes with irregularly distributed dark blotches (yellow

Table I. Morphological measurements of Diasporus citrinobapheus in comparison with other described species of the genus from western Panama and southern Costa Rica (mean±SD, min-max). See Materials and Methods for abbreviations.

Character	D. citrinobapheus	heus	D. diastema		D. hylaeformis	vis	D. ventrimaculatus	culatus	D. vocator		D. tigrillo
	male (n=6)	female (n=1)	male (n=20)	female (n=22)	male (n=9)	female (n=5)	male (n=6)	female (n=2)	male (n=4)	female (n=6)	male (n=2)
SVL (mm)	18.7 ± 0.63	21.8	18.7±1.62 (15.9–22.1)	18.7±2.58 (15.0–23.5)	19.1 ± 1.30 (16.9–20.9)	21.2 ± 0.97 (19.2–21.7)	21.8±1.2 (20.2–23.5)	23.9±0.8	15.3±2.18	14.7±2.18	16.8
DW/LF III	0.23±0.03	0.23		0.32 ± 0.07 $(0.21-0.44)$	0.31 ± 0.03 $(0.27-0.36)$	0.32 ± 0.02 $(0.29-0.34)$			0.26 ± 0.06 $(0.19-0.32)$		
DW/LT IV	$0.14\pm0.03 \\ (0.11-0.18)$	0.17	0.23 ± 0.05 $(0.15-0.32)$	0.22 ± 0.05 $(0.11-0.29)$	0.22 ± 0.03 (0.18-0.26)	0.22 ± 0.02 (0.20-0.24)	1	1	0.17 ± 0.03 (0.13-0.19)	0.17 ± 0.03 (0.14-0.19)	
HL/SVL	$0.41\pm0.01 \\ (0.39-0.44)$	0.40	0.39 ± 0.02 (0.35-0.44)	0.41 ± 0.02 (0.36-0.44)	0.39 ± 0.02 $(0.35-0.43)$	0.39 ± 0.02 $(0.37-0.42)$	0.33	0.35	0.38 ± 0.02 (0.35-0.41)	0.38 ± 0.02 (0.35-0.42)	0.39 (0.38–0.40)
HW/SVL	0.37 ± 0.01 (0.35-0.38)	0.36	0.36 ± 0.02 $(0.33-0.39)$	0.36 ± 0.02 $(0.32-0.39)$	0.37 ± 0.01 (0.35-0.39)	0.36±0.02 (0.35–0.40)	0.39	0.40	0.34 ± 0.02 $(0.31-0.36)$	0.34 ± 0.02 $(0.31-0.36)$	0.36 (0.34–0.37)
HW/HL	0.91±0.03 (0.88–0.97)	06.0	0.91 ± 0.06 (0.79–1.01)	0.90 ± 0.06 (0.78–1.04)	0.94 ± 0.05 $(0.85-1.00)$	0.92 ± 0.04 (0.85-0.96)	1.15	1.14	0.89 ± 0.08 (0.79 ± 0.96)	0.89 ± 0.08 (0.86-0.95)	0.92 (0.85–0.99)
TL/SVL	$0.41 \pm 0.01 \\ (0.40 - 0.42)$	0.42	0.40 ± 0.04 $(0.35-0.51)$	0.42 ± 0.05 $(0.36-0.56)$	0.39 ± 0.01 (0.37–0.42)	0.39 ± 0.05 $(0.35-0.47)$	0.50	0.51	0.40 ± 0.02 $(0.38-0.43)$	0.38 ± 0.02 (0.36-0.42)	0.48 (0.46–0.50)
EL/IOD	$0.98\pm0.12 \\ (0.83-1.12)$	0.94	1.04 ± 0.10 $(0.89-1.24)$	1.12 ± 0.18 $(0.89-1.62)$	1.01 ± 0.12 (0.88–1.24)	1.07 ± 0.10 (0.88–1.19)	98.0	1.00	1.07 ± 0.12 $(0.95-1.24)$	1.43 ± 0.12 (1.25–1.59)	
ED/HL	0.32 ± 0.03 $(0.28 - 0.36)$	0.32	0.29 ± 0.04 $(0.22-0.35)$	0.29 ± 0.04 (0.21–0.37)	0.30 ± 0.03 $(0.27-0.35)$	0.28 ± 0.03 $(0.22-0.30)$	0.37	0.39	0.33 ± 0.01 $(0.32-035)$	$\begin{array}{c c} 0.34\pm0.01 & 0.32 \\ (0.33-0.37) & (0.28-0.35) \end{array}$	0.32 (0.28–0.35)
TD/ED	0.39 ± 0.07 $(0.32-0.45)$	0.32	0.38 ± 0.09 (0.27–0.65)	0.37 ± 0.08 (0.19-0.52)	0.42 ± 0.07 (0.30–0.52)	0.45 ± 0.03 $(0.42-0.50)$	0.48	0.47	0.36 ± 0.07 $(0.30-0.44)$	0.43±0.07 0.55 (0.33–0.50) (0.54–0.57)	0.55 (0.54–0.57)
ED/SVL	0.13 ± 0.01 (0.11-0.15)	0.13	0.11 ± 0.20 (0.08-0.13)	0.11±0.20 0.12±0.13 0.12±0.01 (0.08-0.13) (0.09-0.15) (0.10-0.14)	0.12 ± 0.01 (0.10–0.14)	0.11 ± 0.01 (0.09–0.12)	0.12	0.13	0.13 ± 0.01 (0.12-0.13)	0.13±0.01 0.13±0.01 0.12 (0.12-0.13) (0.12-0.14) (0.11-0.13)	0.12 (0.11–0.13)

to orange dorsal coloration with dark brown spots confined to the pustules); ventral surfaces almost colorless and transparent, in some individuals with a fine dirty white speckling, except male vocal sac that is suffused with yellow (undersurfaces, including venter, yellow); coloration in preservative grayish-white with only a suggestion of yellow (brownish-ocher with dark brown dots; see comments in Discussion section for the usage of different preservation methods). Furthermore, D. citrinobapheus superficially resembles the South American D. gularis from western Ecuador and western Colombia in coloration (see photo in Lynch 2001, page 295 Fig. 7). Diasporus gularis has been described comprehensively by Lynch and Duellman (1997). According to them, adult D. gularis are larger (SVL in males 20.2-21.6 mm, in females 23.3-24.8 mm) than D. citrinobapheus (males 17.3–19.7 mm, single known female 21.8 mm). Moreover, D. gularis shows basal webbings between toes and some specimens have papillae at the apex of the disk pad on some toes, whereas there are no such papillae, and no webbing between toes of *D. citrinobapheus*. The posterior surfaces of thighs are brown in D. gularis, but yellow to orange in D. citrinobapheus. Moreover, the choanae are long, oval, and not concealed by the palatal shelf of the maxillary arch in *D. gularis*, whereas they are round, orientated extremely laterally on palate, and partially concealed by the palatal shelf of the maxillary arch in *D. citrinobapheus*.

Description of the holotype. An adult male; measurements (in mm): SVL 18.4, LF III 2.4, LT IV 4.2, DWF III 0.6, DWT IV 0.5, HL 7.2, HW 7.0, TL 7.8, EL 2.6, IOD 2.9, TD 0.8, ED 2.4; dorsal skin smooth; venter coarsely areolate; no discoidal fold; upper eyelid smooth; snout subovoid in dorsal outline and subacuminate in profile; nostrils weakly protuberant, directed dorsolaterally; head slightly longer than wide, width 97% of length; HW 38% of SVL; canthus rostralis indistinct; ED 36% of HL and 13% of SVL; EL 90% of IOD; TD 33% of ED (Fig. 2 A); choanae round, orientated extremely laterally on palate, partially concealed by palatal shelf of maxillary arch; elliptical vomerine odonthophores, posteromedian to choanae, which are widely separated from each other, with four rows of weakly developed, short teeth; legs short in relation to body; TL 42% of SVL; relative finger length: I<II=IV<III; all fingers with disks, slightly wider than digits, on Fingers II-IV wider than on Finger I; relative toe length: I<II<III<V<IV, Toe V much longer than toe III; tip of Toe V extending to distal subarticular tubercle on Toe IV; tip of Toe III extending to penultimate subarticular tubercle on Toe IV; disks on Toes III-V larger than on I-II; disk covers spadate, lacking papillae; no supernumerary tubercles (Figs 2 B,C).

Etymology. The specific name *citrinobapheus* is a noun in apposition and is derived from the Greek words *citrinos* (citrin-yellow) and *bapheus* (dyer) referring to the yellow body color that dyes one's fingers yellowish when the frog is handled. Although we could observe this phenomenon in a few other species of *Diasporus* too, it is notably evident in the new species.

Coloration in life. All examined specimens show shades of bright yellow and orange dorsally; some have dark grayish and/or whitish-grayish spots (Fig. 3). Ventral surfaces are almost achlorophyllaceous and transparent apart from the yellow male vocal sac.

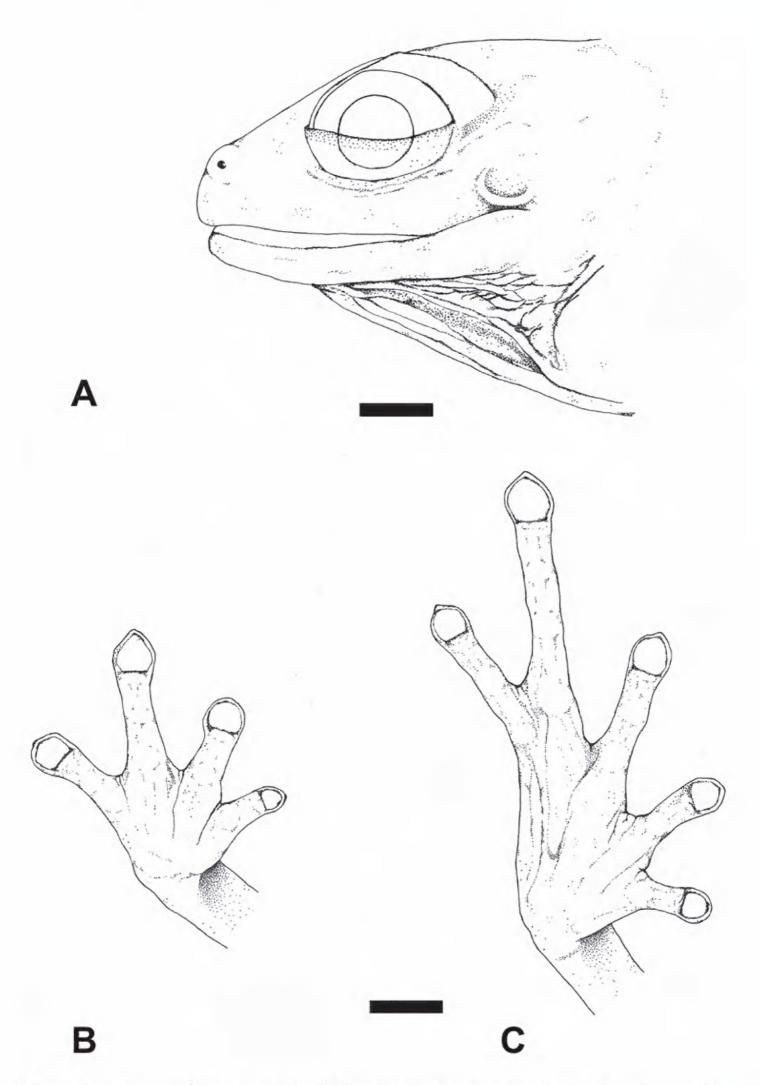


Figure 2. Holotype of *Diasporus citrinobapheus* (SMF 89814, adult male): **A** Lateral view of head **B** Ventral view of right foot. Scale bars = 1 mm.

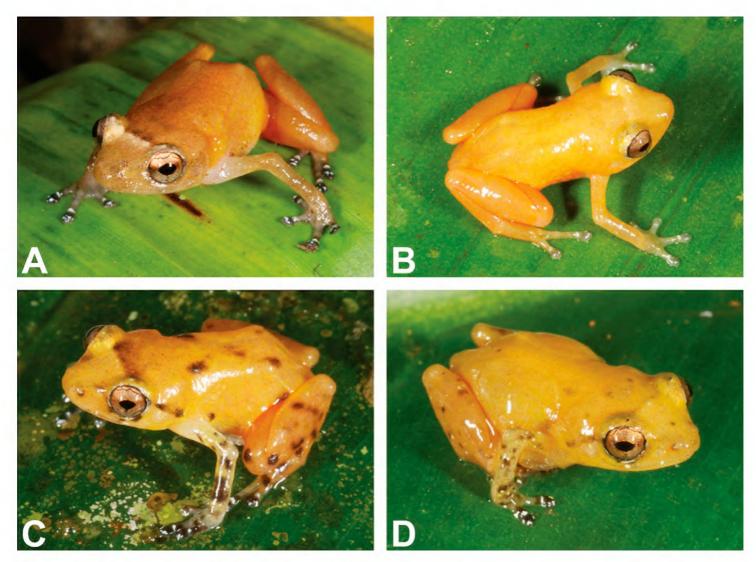


Figure 3. Variation in coloration pattern in life of *Diasporus citrinobapheus* from different localities: **A** Female SMF 89820 from Cerro Negro, Parque Nacional de Santa Fé (Veraguas, Panama) with dirty orange coloration **B** Male SMF 89816 from type locality Paredón (Comarca Ngöbe-Buglé, Panama) with immaculate yellow coloration **C** Male MHCH 2372 from Willie Mazú (Comarca Ngöbe-Buglé, Panama) with intense mottling **D** Male SMF 89817 from Willie Mazú (Comarca Ngöbe-Buglé, Panama) with intermediate mottling.

MHCH 2372 (Fig. 3 C): Dorsal ground color Orange Yellow (18); posterior and anterior surfaces of thighs Chrome Orange (16); Raw Umber (23) interorbital and postocular stripes formed by very fine mottling; dorsum with five Dark Grayish Brown (20) blotches, forming a pattern like the five dots on a dice; scattered Dark Grayish Brown (20) blotches on dorsal surfaces of limbs; disk covers Blackish Neutral Gray (82), with white rings at the base; ventral surface of hind limbs Chrome Orange (16); ventral surface of body transparent with dirty white mottling; vocal sac white with a suggestion of Spectrum Yellow (55).

SMF 89820 (Fig. 3 A): In the only female, coloration in life has been recorded as follows: Dorsal surface Yellow Ocher (123 C); a Chamois (123 D) interorbital bar; anterior and posterior surfaces of thighs Chrome Orange (16); venter almost transparent; upper surfaces of disks Sepia (119) with dirty white spots and a dirty white ring around base; gular region Smoke Gray (44).

Coloration in preservative (70% alcohol). In preservation the bright yellow and orange colors fade rapidly to a pale grayish yellow (Fig. 1 B) with scattered dark gray-

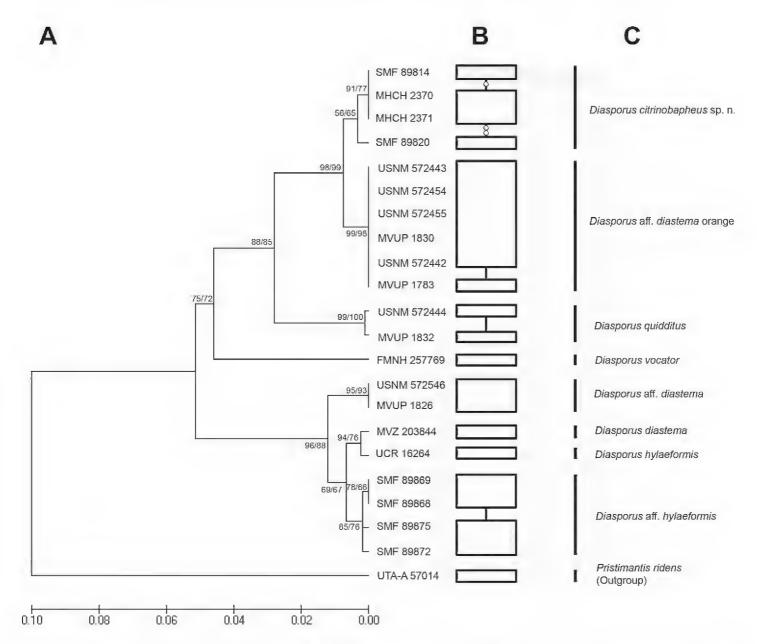


Figure 4. Results of 16S mtDNA analysis. **A** Consensus tree from Maximum Likelihood analysis. Scale bar refers to substitutions per site. Bootstrap support values before the slash correspond to Maximum Likelihood analysis, those after the slash to the Maximum Parsimony consensus tree of exactly the same topology. Numbers behind branches refer to respective museum numbers **B** Parsimony network derived from the same alignment, with each node representing a unique haplotype separated by one substitutional step from its nearest neighbor. Rectangles are haplotypes of analyzed sequences, circles are haplotypes missing in our sample **C** Tentative taxonomic implication. Bar breaks indicate assumed species boundaries. Names refer to morphological determination or GenBank taxonomic identity.

ish blotches in some individuals. Legs pale orange; vocal sac pale yellow in males; gular area in females pale gray; tips of digits dark grayish black. Dark grayish black eyeballs shining through skin when head is viewed dorsally.

Variation. Compared to other species of this genus, the individuals of *Diasporus citrinobapheus* available to us exhibit only little variation in their coloration (Fig. 3). All show a yellow to orange dorsal ground color in life. This can either appear bright and clear or somewhat dirty, depending on the pigment translocation within the melanophores in the frog's skin. In some individuals, higher concentrations of melanophores in certain areas of the dorsum form dark blotches or stripes. This is especially the case in the two specimens from Willie Mazú (Figs 3 C, D). The most frequent pattern of

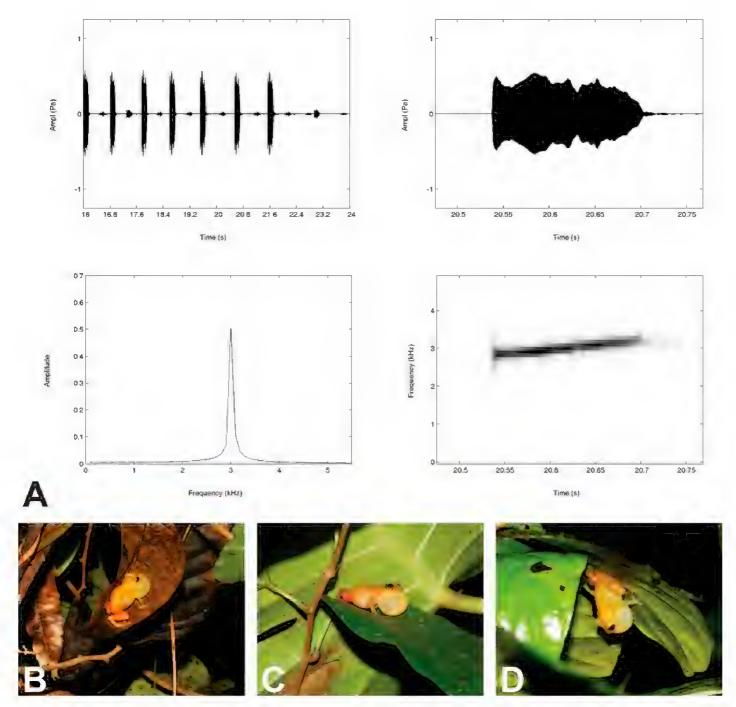


Figure 5. A Visualizations of an advertisement call (Hanning window function, FFT 512, 0.8 overlap) of *Diasporus citrinobapheus* (holotype, SMF 89814) recorded in Paredón, Comarca Ngöbe-Buglé, Panama, at 24.5°C air temperature and 95.3% relative humidity. Clockwise from top left: Oscillogram of a call group; Oscillogram of the penultimate call in the shown call group; Power spectrum showing the dominant frequency of the penultimate call in the shown call group; Spectrogram of the penultimate call in the shown call group **B–D** Different call positions of male *D. citrinobapheus*: **B** Male holotype (SMF 89814) from Paredón calling on dead leaves in dense vegetation about 2 meters above ground level; **C** Male paratype (MHCH 2371) from Paredón on green leaf about 3 m above ground level **D** Male specimen (SMF 89817) from Willie Mazú referred to as *D. citrinobapheus* calling from an elevated position on the underside of a leaf.

this type is an interorbital bar, which in most cases is darker than ground color along the anterior edge of the bar and lighter than ground color along the posterior edge. In addition, some individuals show dark brown blotches on the limbs and less frequently also on the dorsum. Most individuals show additional small whitish spots, in particular under and around the eyes, as well as scattered across the forelimbs. In the male SMF 89816 from the type locality (Fig. 3 B) the dark and white markings on and around the disk covers are not as pronouncedly contrasting as in the other individuals examined.

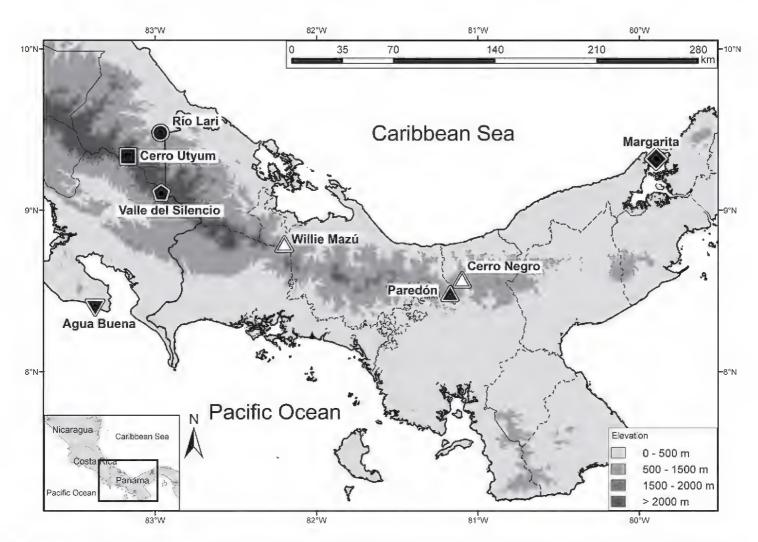


Figure 6. Distribution map of *Diasporus citrinobapheus* and type localities of other species in the genus in Panama and Costa Rica. **Solid triangle:** Paredón, Comarca Ngöbe-Buglé, type locality of *D. citrinobapheus*. **Hollow triangles:** Additional collection sites of *D. citrinobapheus*: Private Reserve Willie Mazú in the west, and Cerro Negro (Parque Nacional Santa Fé, Veraguas) in the east. **Inverted triangle:** Agua Buena, Puntarenas, Costa Rica, type locality of *D. vocator*. **Pentagon:** Valle del Silencio, at the provincial boarder between Puntarenas and Limón, Costa Rica, type locality of *D. ventrimaculatus*. **Square:** Cerro Utyum, Limón, Costa Rica, type locality of *D. hylaeformis*. **Circle:** Río Lari, Limón, Costa Rica, type locality of *D. tigrillo*. **Diamond:** Margarita, Colón, Panama, type locality of *D. diastema*. Dashed lines represent provincial borders. Solid lines represent coast line and national border.

Molecular genetics. The distinctiveness of *Diasporus citrinobapheus* is supported by the analysis of the 16S mitochondrial rRNA gene (Fig. 4). The four individuals we examined form a distinct cluster that appears separated from the other members for which 16S sequences are available. The mean genetic distance among the four specimens of *D. citrinobapheus* is 0.3%. In our consensus tree (Fig. 4 A) it appears to be most closely related to the candidate species *D.* aff. *diastema* from El Copé, from which it is separated by a mean genetic distance of 1.8%. In the haplotype network analysis (Fig. 4 B) both clades form unconnected subnetworks, indicating a differentiation at species level (Fig. 4 C). The mean genetic distance to the next closest relative *D. quidditus* is 6.6% for *D. citrinobapheus* and 7% for *D. aff. diastema*.

Vocalization of holotype. We recorded a 3 min, 43.5 seconds portion of the advertisement call of the holotype that yielded a total of 63 calls. An exemplary visualization of the call structure is given in Fig. 5 A. Relative humidity during recording was 95.3%

at an air temperature of 24.5 °C. As in other members of the genus, the call consists of a single note, even though calls sound like a "whistle," rather than the typical "tink" usually emitted by members of the genus *Diasporus* (Savage 2002; Chaves et al. 2009). The 63 recorded calls are organized in five call groups of 8–17 calls per group (12.8 \pm 3.2). A call group lasts 19.8–34.1 s (25.0 \pm 5.7). Intervals between call groups range from 15.7–33.2 s (21.6 \pm 8.0) and intervals between calls within a call group range from 0.57–5.77 s (1.93 \pm 1.2). Call group rate is 1.34 call groups per minute; call rate within a call group varies from 23.4–40.8 calls per minute (32.0 \pm 6.3). Call duration varies from 0.13–0.18 s (0.16 \pm 0.01). There is a rather weak frequency modulation of 190–470 Hz (370 \pm 65). The spectrum of frequencies within a call range from a mean minimum of 2890 \pm 44 Hz to a mean maximum of 3260 \pm 44 Hz. Fundamental and dominant frequencies are identical at about 2950 Hz. The dominant frequency, as the frequency with the greatest energy in the signal, is reached about 0.05 s after initiation of the call.

Vocalizations of paratypes and referred specimens. In addition to the holotype, we recorded and analyzed the advertisement calls of two paratypes (SMF 89816, MHCH 2371) and one referred specimen (SMF 89817). Summing up, the advertisement call of *Diasporus citrinobapheus* sounds like a whistle, is organized in call groups, has a call duration of 0.14–0.16 s in average and a dominant frequency of 2860–3040 Hz (see all parameters in Table 2). While the paratypes vary only little in call parameters, SMF 89817 shows obvious differences regarding call duration, call interval, and call rate (see Discussion section for details).

Geographical distribution and natural history notes. So far, *Diasporus citrinobapheus* has been found on the Caribbean slopes of the western Serranía de Tabasará and on both Pacific and Caribbean slopes of the eastern Serranía de Tabasará (Fig. 6) at intermediate elevations from 680 to 790 m.a.s.l. Males call from very dense vegetation and are difficult to spot. They are almost only detectable by following their characteristic vocalization. Vocal activity is highest just after dusk and finally stops when it

Table 2. Variation in advertisement call parameters in four male specimens referred to as *Diasporus citrinobapheus* (mean±SD, min–max).

	SMF 89814	SMF 89816	MHCH 2371	SMF 89817
Temperature / RH during recording	24.5° C/95.3 %	24.3° C/93.5 %	24.6° C/93.6 %	21.8° C/100 %
Total recording time (min)	3.73	1.35	1.66	3.03
Number of call groups recorded	5	2	1	4
Number of calls recorded	63	26	11	68
Call group rate (call groups / min)	1.34	1.48	0.6	1.32
Call group duration (s)	25.0±5.7 19.8–34.1	23.0±9.5 16.3–29.7	19.0	20.6±8.5 12.5–28.5
Calls per group	12.8±3.2 8–17	11–15	11	16.6±5.7 10–22

	SMF 89814	SMF 89816	MHCH 2371	SMF 89817
Call group interval (s)	21.6±8.0 15.7–33.2	26.84	>78	25±16.7 10.9–43.5
Call rate over entire recording (calls/min)	16.9	19.3	6.63	22.4
Call rate within a call group (calls/min)	32±6.3 23.4–40.8	35.4±7.1 30.3–40.4	35	50±6.8 44.2–60
Call duration (s)	0.157±0.01 0.126–0.178	0.162±0.01 0.143–0.174	0.156±0.003 0.151 – 0.162	0.141±0.01 0.114–0.167
Call interval (s)	1.93±1.2 0.57–5.77	1.74±1.4 0.63–3.77	1.71±0.75 0.85 – 2.85	1.15±0.49 0.55–2.58
Dominant frequency (Hz)	2953±0	3010±75 2859–3140	2859±0	2965±32 2953–3046
Minimum frequency (Hz)	2889±44 2859–2953	2776±31 2765–2859	2671±0	2939±33 2895–2953
Maximum frequency (Hz)	3257±44 3140–3328	3184±61 3064–3234	3029±38 2953–3046	3290±74 3140–3421
Frequency modulation (Hz)	367 188–469	407 281–468	358 281–375	351 281–375

becomes dark. Calling height ranges from near ground level up to three meters above ground. Calling position can be either on the upper side of a leaf or on its underside (Figs 4 B–D). The only female (SMF 89820) was found at daytime (15:00 h) inside an involute, young plantain leaf that apparently served as a daytime hiding place. The species does not seem to be limited to mature forest, but is also found in secondary growth and plantations. However, it appears to avoid open habitats like pasture land.

Discussion

Diasporus citrinobapheus is easily distinguishable from all other known frogs of the genus in Lower Central America by its bright yellow to orange coloration. The only described species of the genus that somewhat resemble the new species in coloration are *D. gularis* from Colombia and Ecuador and *D. tigrillo* from the Caribbean slopes of the Costa Rican part of the Serranía de Talamanca. The latter species is known only from two specimens, both collected in 1964 at a single locality and there are no photographs of the species in life, tissue samples, or call recordings available to clarify the systematic relationships of this species. The different ground coloration in preservative between *D. tigrillo* and *D. citrinobapheus* is certainly due to different preservation techniques, because the fixation process in 10% formalin darkens the complete specimen. However, this does not influence the general color pattern, so we treat the dark brown spots on the dorsum of *D. tigrillo* as a diagnostic feature to differentiate between *D. tigrillo* and *D. citrinobapheus*. Additional material is required, preferably from near the type locality of *D. tigrillo* to conduct further studies. In contrast, *D. gularis* is known

from a number of specimens from Colombia and Ecuador. However, the presence and development of papillae at the apex of the pad on the underside of the disk cover, one of the main characters that has been used to distinguish this species from its congeners, is a controversial issue. Lynch (1976, page 12, Fig. 3 B) provided a drawing of the left hand of a specimen (LACM 73239) from Chocó, Colombia, that shows long papillae. In a later work, the same author presented drawings of finger disk pads of two D. gularis (ICN 45168, 45171) from Valle del Cauca, Colombia, that show knobbed disk covers (Lynch 2001, Figs 2 A, B), but he also noted that there are specimens lacking this character. He further considered the presence or absence of this knob at the underside of the disk cover might be due to preparation technique. Lynch and Duellman (1997) noted that the holotype of *D. gularis* from Ecuador does not have papillae at the tip of any digit, while they stated that specimens from southern Colombia and Ecuador have a rounded knob at the apex of the pad on the underside of the disk cover of toe II-IV. In Chocoan Colombia, specimens referred to as D. gularis have larger papillae. Depending on their diagnosis, Lynch and Duellman (1997) argued that several species might currently be referred to D. gularis, three in western Colombia and only one in Ecuador, where the type locality is. Based on our genetic analysis, D. citrinobapheus is closely related to, and may even be conspecific with, the candidate species D. aff. diastema from El Copé (Crawford et al. 2010a). Albeit the comparably small p-distance, the haplotype network analysis yields a separate network for each of the two clades, supporting the assumption of two distinct species. However, genetic evidence revealed only from mitochondrial markers alone is not strong enough to support either one or the other hypothesis (Vences et al. 2005). Further integrative taxonomic studies, including morphology, bioacoustics, and nuclear genes are needed to clarify this matter.

Besides various records of other amphibians and reptiles, we found no additional species of the genus *Diasporus* at the type locality. At Willie Mazú, a locality approximately 120 km NW of the type locality of *D. citrinobapheus*, we collected a single specimen of *Diasporus* that we refer to *D. vocator* based on size, coloration, disk shape, and male advertisement call. At Cerro Negro, *D. citrinobapheus* occurs sympatrically with *D. diastema*. Based on our current concept of its distribution, the possibility remains that also *D. vocator* occurs at this locality, although the species has not been recorded from this site.

The eponymous, readily soluble yellow coloration of *Diasporus citrinobapheus* lead us to the assumption that this might serve a defensive function against predators. On this account, an alcohol extraction was analyzed for alkaloids, but no active substances were found. Probably, the yellow pigment is just highly soluble and therefore easily washed out. Nevertheless, one could speculate that it has a bitter or otherwise unpalatable taste that might deter certain predators.

Various studies have shown that the advertisement call represents a premating isolating mechanism in anurans (e.g., Duellman and Trueb 1986), which makes it a valuable tool in taxonomy. Having in mind that there are great morphological overlaps between members of the genus *Diasporus*, analyses of vocalizations might form the most powerful taxonomic approach to decipher its species diversity. Unfortunately,

the calls of most species have never been formally described. Fouquette (1960) was the first to describe the call of *D. diastema* from the Panama Canal area, about 10 km northwest of Panama City, not far from the type locality (Cope 1876; Dunn 1942). Later, Wilczynski and Brenowitz (1988) presented another call description based on calls recorded in the surroundings of Gamboa in Central Panama, about 24 km NNW of Panama City, and even closer to the type locality. Interestingly, the call descriptions of Fouquette (1960) and Wilczynski and Brenowitz (1988) are incongruent in terms of call duration, frequency range, and dominant frequency, rendering it possible that different species were recorded. Unfortunately, none of these papers cited any voucher specimens, so it is impossible to determine which species they actually recorded. The most recent contribution on vocalizations of *D. diastema* is that of Ibáñez et al. (1999), also from the environs of the Panama Canal. They provided a rough sonogram, but did not give any numerical values. The dominant frequency in all three papers is roughly described as 3000–4000 Hz, thus considerably higher than in D. citrinobapheus. Furthermore, all three papers (Fouquette 1960; Wilczynski and Brenowitz 1988; Ibáñez et al. 1999) present sonograms, which show an obvious frequency modulation expressed by a rapid rise of frequency over time with approximately 1000 Hz difference between beginning and end of the call. In contrast, the call of *D. citrinobapheus* is characterized by only a moderate frequency rise over time, on average 350-400 Hz. Confusing are the data of call duration provided by Fouquette (1960) and Wilczynski and Brenowitz (1988), respectively. Fouquette (1960) reports on mean call duration of 0.2 s for D. diastema. Yet, in the accompanying sonogram (Fig. 2 A in Fouquette 1960), the call seems to be only slightly longer than 0.1 s. Wilczynski and Brenowitz (1988) even mentioned a call duration of more than 0.3 s, but in the accompanying oscillogram (Fig. 1 B in Wilczynski and Brenowitz 1988), the call does not exceed 0.1 s on the time axis. Although difficult to assess precisely, the duration of the call pictured in the sonogram provided by Ibáñez et al. (1999) is clearly shorter than 0.2 s. Furthermore, Ibáñez et al. (1999) present a sonogram of *D. vocator*, recorded also in the Canal Zone. According to their analysis, the call of *D. vocator* has a frequency range between 6000 and 7000 Hz, is very short, and shows a strong frequency modulation, thus being very different from the calls of D. diastema and D. citrinobapheus. However, the type locality of *D. vocator* is Agua Buena in the Peninsula de Osa, Costa Rica. Thus, it is advisable to record comparative call material from the type locality for future analyses. The most recent contribution on *Diasporus* vocalizations was made by Chaves et al. (2009) in the course of the original description of *D. ventrimaculatus*. This species' voice differs in all standard parameters from that of *D. citrinobapheus*, as it has much shorter call durations of about 0.08 s, a low dominant frequency of about 2550 Hz, and a lower frequency range between 2140 and 2995 Hz. Furthermore, the dominant frequency is reached at the very beginning at the call. The same authors presented a preliminary analysis of calls emitted by specimens assigned to D. hylaeformis. According to this analysis, call duration in D. hylaeformis is on average 0.214 s, while it resembles D. ventrimaculatus in spectral parameters. Regarding the vocalizations of D. tigrillo, the least known species in the genus, only a field note citation appearing in the original

description describes it as "similar to the dink dink of *Eleutherodactylus [Diasporus]* diastema" (Savage 1997).

Nevertheless, there is also an intraspecific variation among calls of specimens referred to *Diasporus citrinobapheus*. The call of the single male recorded at Willie Mazú (SMF 89817) differs from the calls of the members of the type series in temporal parameters, such as shorter call duration and call interval that result in a higher call rate. These differences are minor, but lead us to not include specimens from localities other than the type locality in the type series. However, various studies have shown that call parameter variation is linked to ambient temperature (e.g., Zweifel 1959; Schneider 1977; Gerhardt 1978). According to these studies, call duration and call interval are negatively correlated with temperature, which in turn leads to an increased call rate at higher temperatures. As SMF 89817 was recorded at lower temperatures than for the other three specimens, one would expect the opposite pattern. Nevertheless, these studies used data from many individuals, built scatter diagrams of parameters against temperature and fitted least-squares regression, and there are always outliers that do not follow the general trend. In our case, individual differences may be stronger than temperature-related ones, but this assumption needs further research to be reliably assessed.

Apart from morphology which apparently is not the best tool to identify species of *Diasporus*, neither DNA nor bioacoustics, both of paramount importance for contemporary anuran taxonomy, have been consistently analyzed among geographically and taxonomically wide-ranging samples. While the Panamanian and Costa Rican 16S barcodes compared in this study reveal the existence of more infrageneric lineages than names are available, the doubtless assignation of a given *Diasporus* "aff. *hylaeformis*" or "aff. *diastema*" is likely to be highly challenging if one is to rely on the existing treatments, which mostly provide only partial or even contradicting information. In conclusion, the complex and cryptic diversity within the genus *Diasporus* requires a thorough revision of as many "quality vouchers" (collected specimens associated with both well-preserved tissue samples and call recordings) from as many localities throughout the generic range as possible.

Key to the species of Diasporus in Central America

la	Disk covers lanceolate or papillate2
1b	Disk covers palmate or spadate
2a	Dorsum shagreened; fingers without thick lateral fringes; Toe V not partially
	fused with Toe IV; SVL of adult males 14.0–16.0 mm, of adult females 16.5–
	18.0 mm
2b	Dorsum with scattered low warts; fingers with thick lateral fringes; Toe V par-
	tially fused with Toe IV; SVL of adult males 10.9-14.8 mm, of adult females
	13.2–16.9 mm
3a	Fingers II and III with palmate disk covers and broadened, non-triangular
	disk pads; adults with vomerine teeth

3b	Fingers II and III with spadate disk covers and triangular disk pads; adults
	with or without vomerine teeth4
4a	Venter in most individuals with distinct black and white blotches; dorsum
	and dorsal surfaces of arms and legs in some individuals bright red in life
	Diasporus ventrimaculatus
4b	Venter patternless or with a few small black dots; dorsum and dorsal surfaces
	of arms and legs brown, cream, or yellow in life5
5a	Posterior surface of thigh pigmented (brownish, often suffused with red in
	life); overall dorsal coloration bright cream, grayish or reddish brown in life;
	adults without vomerine teeth
5b	Posterior surface of thigh unpigmented (yellow in life); overall dorsal colora-
	tion bright yellow to orange in life; adults with vomerine teeth6
6a	Dorsum with scattered low pustules; ratio tympanum length / eye length
	0.54-0.57; distal subarticular tubercle on Fingers and Toes I weakly bifid;
	dorsum yellow to orange with dark brown spots confined to pustules; SVL of
	adult males 16.0–17.5 mm
6b	Dorsum smooth; ratio tympanum length / eye length 0.32-0.45; distal sub-
	articular tubercle on Fingers and Toes I flat and rounded; dorsum uniformly
	bright yellow to orange, sometimes with irregularly distributed dark blotches;
	SVL of adult males 17.3–19.7 mm

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References

- Bee MA (2004) Equipment Review: Sound Ruler Acoustical Analysis: a free, open code, multi-plat-form sound analysis and graphing package. Bioacoustics 14: 171–178. http://soundruler.source-forge.net/main/docs/sruler_review_2004_bee.pdf, doi: 10.1080/09524622.2004.9753520
- Bolaños F, Wake DB (2009) Two new species of montane web-footed salamanders (Plethodontidae: *Bolitoglossa*) from the Costa Rica-Panama border region. Zootaxa 1981: 57–68. http://ib.berkeley.edu/labs/wake/349_Bol_Wake_Zoot_09-1.pdf
- Chaves G, García-Rodríguez A, Mora A, Leal A (2009) A new species of dink frog (Anura: Eleutherodactylidae: *Diasporus*) from Cordillera de Talamanca, Costa Rica. Zootaxa 2088: 1–14. http://www.mendeley.com/research/a-new-species-of-dink-frog-anura-eleuthero-dactylidae-diasporus-from-cordillera-de-talamanca-costa-rica
- Clement M, Posada D, Crandall K (2000) TCS: a computer program to estimate gene genealogies. Molecular Ecology Notes 9: 1657–1660.
- Cope ED (1876) On the Batrachia and Reptilia of Costa Rica. Journal of the Academy of Natural Sciences of Philadelphia, Series 2, 8: 93–154. http://www.iucn-tftsg.org/wp-content/uploads/file/Articles/Cope_1876.pdf
- Crawford AJ, Lips KR, Bermingham E (2010a) Epidemic disease decimates amphibian abundance, species diversity, and evolutionary history in the highlands of Central Panama. Proceedings of the National Academy of Science 107 (31):13777–13782. doi: 10.1073/pnas.0914115107
- Crawford AJ, Ryan MJ, Jaramillo CA (2010b) A new species of *Pristimantis* (Anura: Strabomantidae) from the Pacific coast of the Darien Province, Panama, with a molecular analysis of its phylogenetic position. Herpetologica 66: 171–185. doi: 10.1655/09-018R1.1
- Drummond AJ, Ashton B, Cheung M, Heled J, Kearse M, Moir R, Stones-Havas S, Thierer T, Wilson A (2010) Geneious v.4.8.5. www.geneious.com (accessed 27 February 2012).
- Duellman WE, Trueb L (1986) Biology of amphibians. The Johns Hopkins University Press, London, 696 pp.
- Dunn ER (1942) A new species of frog (*Eleutherodactylus*) from Costa Rica. Notulae Naturae of the Academy of Natural Sciences of Philadelphia 104: 1–2.
- Fouquette MJ JR (1960). Call structure in frogs of the family Leptodactylidae. The Texas Journal of Science 12 (3): 201–215.
- Frost DR (2011) Amphibian Species of the World: an Online Reference. Version 5.5 (31 January, 2011). American Museum of Natural History, New York, USA. http://research.amnh.org/vz/herpetology/amphibia/ (accessed 1 November 2011).
- Gerhardt HC (1978) Temperature coupling in the communication system of the Gray Tree Frog, *Hyla versicolor*. Science 199: 992–994. doi: 10.1126/science.199.4332.992
- Gridi-Papp M (Ed.) (2003–2007) SoundRuler: Acoustic Analysis for Research and Teaching. http://soundruler.sourceforge.net (accessed 1 March 2011).
- Hart MW, Sunday J (2007) Things fall apart: biological species form unconnected parsimony networks. Biology Letters 3: 509–512. doi: 10.1098/rsbl.2007.0307
- Hedges SB, Duellman WE, Heinicke MP (2008) New World direct-developing frogs (Anura: Terrarana): Molecular phylogeny, classification, biogeography, and conservation Zootaxa 1737: 1–182. http://www.mapress.com/zootaxa/2008/2/zt01737p182.pdf

- Ibáñez R, Solís F, Jaramillo C, Rand S (2001) An overview of the herpetology of Panama. In: Johnson JD, Webb RG, Flores-Villela OA (Eds) Mesoamerican Herpetology: Systematics, Zoogeography, and Conservation. Centennial Museum Special Publications No. 1, The University of Texas, El Paso, 159–170.
- Ibáñez R, Rand AS, Jaramillo CL (1999) Los anfibios del Monumento Natural Barro Colorado, Parque Nacional Soberania y áreas adyacentes/The amphibians of Barro Colorado Nature Monument, Soberania National Park and adjacent areas. Editorial Mizrachi and Pujol S.A., Panamá, 187 pp.
- IUCN (2011) IUCN Red List of Threatened Species. Version 2011.1. www.iucnredlist.org [accessed 1 November 2011]
- Ivanova NV, De Waard J, Hebert PDN (2006) An inexpensive, automation-friendly protocol for recovering high-quality DNA. Molecular Ecology Notes 6: 998–1002. doi: 10.1111/j.1471-8286.2006.01428.x
- Jaramillo C, Wilson LD, Ibáñez R, Jaramillo F (2010) The herpetofauna of Panama: distribution and conservation. In: Wilson LD, Townsend JH, Johnson JD (Eds) Conservation of Mesoamerican Amphibians and Reptiles, Eagle Mountain Publications LC, Eagle Mountain, Utah, 604–671.
- Köhler G (2001) Anfibios y reptiles de Nicaragua. Herpeton Verlag Elke Köhler, Offenbach, Germany, 208 pp.
- Köhler G (2011) Amphibians of Central America. Herpeton Verlag Elke Köhler, Offenbach, Germany, 378 pp.
- Köhler G, Ponce M, Batista A (2007) A new species of worm salamander (genus *Oedipina*) from Fortuna, western central Panama. Senckenbergiana Biologica 87: 213–217.
- Larkin MA, Blackshields G, Brown NP, Chenna R, McGettigan PA, McWilliam H, Valentin F, Wallace IM, Wilm A, Lopez R, Thompson JD, Gibson TJ, Higgins DG (2007) ClustalW and ClustalX version 2. Bioinformatics 23 (21): 2947–2948. doi: 10.1093/bioinformatics/btm404
- Lynch JD (1976) The species groups of the South American frogs of the genus *Eleutherodactylus* (Leptodactylidae). Occasional papers of the Museum of Natural History The Kansas University, Lawrence 61: 1–24.
- Lynch JD (2001) Three new rainfrogs of the *Eleutherodactylus diastema* group from Colombia and Panama. Revista de la Academia Colombiana de Ciencias Exactas, Físicas y Naturales 25: 287–297.
- Lynch JD, Duellman WE (1997) Frogs of the genus *Eleutherodactylus* in western Ecuador: Systematics, Ecology, and Biogeography. The University of Kansas Natural History Museum, Lawrence, Kansas, USA, 236 pp.
- Mendelson III JR, Mulcahy DG (2010) A new species of toad (Bufonidae: *Incilius*) from central Panama. Zootaxa 2396: 61–68. http://www.mapress.com/zootaxa/2010/f/zt02396p068.pdf
- Mendelson III JR, Savage JM, Griffith E, Ross H, Kubicki B, Gagliardo R (2008) Spectacular new gliding species of *Ecnomiohyla* (Anura: Hylidae) from central Panama. Journal of Herpetology 42: 750–759. doi: 10.1670/08-025R1.1
- Myers CW, Duellman WE (1982) A new species of *Hyla* from Cerro Colorado, and other tree frog records and geographic notes from western Panama. American Museum Novitates 2752: 1–32.

- Ryan MJ, Lips KR, Giermakowski JT (2010a) New species of *Pristimantis* (Anura: Terrarana: Strabomantidae) from lower Central America. Journal of Herpetology 44: 193–200. doi: 10.1670/08-280.1
- Ryan MJ, Savage JM, Lips KR, Giermakowski JT (2010b) A new species of the *Craugastor rugulosus* series (Anura: Craugastoridae) from west-central Panama. Copeia 2010: 405–409. doi: 10.1643/CH-09-154
- Sabaj Pérez MH (Ed.) (2010) Standard symbolic codes for institutional resource collections in herpetology and ichthyology: an Online Reference. Verson 1.5. American Society of Ichthyologists and Herpetologists, Washington, DC., 35 pp. http://www.asih.org/ (accessed 4 October 2010).
- Savage JM (1997) A new species of rainfrog of the *Eleutherodactylus diastema* group from the Alta Talamanca region of Costa Rica. Amphibia-Reptilia 18: 241–247. doi: 10.1163/156853897X00125
- Savage JM (2002) The Amphibians and Reptiles of Costa Rica: A Herpetofauna between two Continents, between two Seas. University of Chicago Press, Chicago, 934 pp. http://www.ingentaconnect.com/content/brill/amre/1997/00000018/0000003/art00003
- Schneider H (1977) Acoustic behavior and physiology of vocalization in the European tree frog, *Hyla arborea* (L.). In: Taylor DH, Guttman SI (Eds) The reproductive biology of Amphibians. Plenum Press Corp., New York, 295–335.
- Smithe FB (1975–1981) Naturalist's color guide. Part I. Color guide. American Museum of Natural History, New York, New York, USA, 182 color swatches.
- Tamura K (1992) Estimation of the number of nucleotide substitutions when there are strong transition-transversion and G + C-content biases. Molecular Biology and Evolution 9: 678–687.
- Tamura K, Peterson D, Peterson N, Stecher G, Nei M, Kumar S (2011) MEGA5: Molecular Evolutionary Genetics Analysis using Maximum Likelihood, Evolutionary Distance, and Maximum Parsimony Methods. Molecular Biology and Evolution 28: 2731–2739. doi: 10.1093/molbev/msr121
- Vences M, Thomas M, Bonett RM, Vieites DR (2005) Deciphering amphibian diversity through DNA barcoding: chances and challenges. Philosophical Transactions of the Royal Society of London Series B 360: 1859–1868. doi: 10.1098/rstb.2005.1717
- Wake DB, Hanken J, Ibáñez R (2005) A new species of big black *Bolitoglossa* (Amphibia: Caudata) from central Panama. Copeia 2005: 223–226. doi: 10.1643/CH-04-082R1
- Wake DB, Savage JM, Hanken J (2007) Montane salamanders from the Costa Rica-Panama border region, with descriptions of two new species of *Bolitoglossa*. Copeia 2007: 556–565. http://www.jstor.org/stable/25140665, doi: 10.1643/0045-8511(2007)2007[556:MSFT-CR]2.0.CO;2
- Wilczynski W, Brenowitz EA (1988) Acoustic cues mediate inter-male spacing in a Neotropical frog. Animal Behavior 36: 1054–1063. doi: 10.1016/S0003-3472(88)80065-4
- Zweifel RG (1959) Effects of temperature on calls of *Bombina variegata*. Copeia 1959: 322–327. http://www.jstor.org/stable/1439890, doi: 10.2307/1439890

Appendix I

Comparative material for morphological examination

Diasporus diastema — Costa Rica: Heredia: Puerto Viejo de Sarapiquí, entrance to La Selva, 30 m: SMF 81812; Rara Avis, 700 m: SMF 81811; Honduras: Gracias a Dios: Region La Mosquitia, Rio Plátano Biosphere Reserve, Raudal Kiplatara, 162 m: SMF 85938; El Ocotilla, 410 m: SMF 85939; Nicaragua: Atlántico Norte: PN Saslaya, 920 m: SMF 82031-82035; Jinotega: Bosawas Biosphere Reserve: SMF 78561; National Park Saslaya, 188 m: SMF 78965; Matagalpa: Selva Negra, 1300 m: SMF 77231, 77235, 78184–78191; Río San Juan: Bartola, 30–70 m: SMF 80977–80979, SMF 79799–79800; Río Sarnoso, 25 m: SMF 79796–79798; Boca de San Carlos, 20 m: SMF 79794–5; ridge near Río Las Cruces, near Caño Negro, 415 m: SMF 83389; Cerro el Bolívar, near Río Machado, 280 m: SMF 83390; Lomas de Tambor, 210 m: SMF 83391; Panama: Bocas del Toro: Bosque Protector Palo Seco, 1148 m: SMF 84997; Archipelago Bocas del Toro, Isla Colón, 30 m: SMF 85068; Chiriquí: El Chorogo, 295 m: SMF 92008; Coclé: Cerro Gaitál, El Valle de Antón, 800 m: SMF 80781; Comarca Kuna Yala: Reserva Natural Nusagandi, 280 m: SMF 81961; Panamá: Canal Zone: SMF 29859, 29874.

Diasporus aff. hylaeformis — Panama: Bocas del Toro: Parque Internacional La Amistad, northern slope of Cerro Pando, 2400 m: SMF 89868, 89869, 89873, 89874, AH 263, AH 266. Chiriquí: Parque Internacional La Amistad, Jurutungo, southern slope of Cerro Pando, 2070–2330 m: SMF 89867, 89872, 89875, 89876, AH 126, 127; Cerro Punta, Las Nubes Ranger Station, 2070 m: SMF 89870, 89871.

Diasporus vocator — **Panama**: Bocas del Toro: Humedal de San San Pond Sak, 5 m: SMF 89865, AH 364; Comarca Kuna Yala: Reserva Natural Nusagandi, 350 m: SMF 81970–81976.

Appendix II

Corresponding information of sequenced *Diasporus* specimens.

Species	Collection number	Field number	GenBank accession no.	Country	Province	Latitude	Longitude
D. citrinobapheus	SMF 89814	AH 449	JQ927333	Panama	Comarca Ngöbe-Buglé	8.485	-81.173
D. citrinobapheus	SMF 89820	AH 211	JQ927334	Panama	Veraguas	8.569	-81.099
D. citrinobapheus	MHCH 2370	AH 450	JQ927335	Panama	Comarca Ngöbe-Buglé	8.485	-81.173
D. citrinobapheus	MHCH 2371	AH 452	JQ927336	Panama	Comarca Ngöbe-Buglé	8.485	-81.173
D. aff. hylaeformis	SMF 89868	AH 267	JQ927337	Panama	Bocas del Toro	8.931	-82.714

Species	Collection number	Field number	GenBank accession no.	Country	Province	Latitude	Longitude
D. aff. hylaeformis	SMF 89869	AH 268	JQ927338	Panama	Bocas del Toro	8.931	-82.714
D. aff. hylaeformis	SMF 89872	AH 124	JQ927339	Panama	Chiriquí	8.911	-82.713
D. aff. hylaeformis	SMF 89875	AH 282	JQ927340	Panama	Chiriquí	8.912	-82.713
D. aff. diastema 'orange'	USNM 572442	KRL 0902	FJ784425	Panama	Coclé	8.667	-80.592
D. aff. diastema 'orange'	USNM 572443	KRL 1181	FJ784484	Panama	Coclé	8.667	-80.592
D. aff. diastema 'orange'	USNM 572454	KRL 0900	FJ784423	Panama	Coclé	8.667	-80.592
D. aff. diastema 'orange'	USNM 572455	KRL 0901	FJ784424	Panama	Coclé	8.667	-80.592
D. aff. diastema 'orange'	MVUP 1783	KRL 0694	FJ784338	Panama	Coclé	8.667	-80.592
D. aff. diastema 'orange'	MVUP 1830	KRL 0840	FJ784395	Panama	Coclé	8.667	-80.592
D. quidditus	USNM 572444	KRL 0647	FJ784326	Panama	Coclé	8.667	-80.592
D. quidditus	MVUP 1832	KRL 0856	FJ784405	Panama	Coclé	8.667	-80.592
D. vocator	FMNH 257769	AJC 0127	JN991419	Costa Rica	Puntarenas	8.79	-82.96
D. aff. diastema	USNM 572546	KRL 0782	FJ784369	Panama	Coclé	8.667	-80.592
D. aff. diastema	MVUP 1826	KRL 0831	FJ784390	Panama	Coclé	8.667	-80.592
D. diastema	MVZ 203844	1999	EU186682	Costa Rica	Cartago	9.75	-83.804
D. hylaeformis	UCR 16264	AJC 0468	JN991418	Costa Rica	Alajuela	10.22	-84.54
Pristimantis ridens	UTA-A 57014	ENS 10722	JN991464	Honduras	Olancho	14.93	-86.14

Appendix III

Audio samples of the advertisement calls of specimens of *Diasporus citrinobapheus*. (doi: 10.3897/zookeys.196.2774.app1) File format: MP3 Audio file (MP3).

Explanation note: MP3 audio samples of specimens of *Diasporus citrinobapheus* recorded at the type locality (Paredón) and Willie Mazú, Serranía de Tabasará, Panama.

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